AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1. (original): A process for producing an optically active 1,4-pentanediol represented by formula (2):

(wherein * represents an asymmetric carbon atom) comprising asymmetrically reducing 5-hydroxy-2-pentanone represented by formula (1):

- 2. (original): The process according to claim 1, wherein 5-hydroxy-2-pentanone represented by said formula (1) is asymmetrically reduced by the action of an enzyme source having the activity of stereoselectively reducing the compound.
- 3. (original): The process according to claim 2, wherein the enzyme source is a cultured product of a microorganism belonging to genus <u>Candida</u>, genus <u>Devosia</u>, genus <u>Rhodococcus</u>, or genus <u>Rhodotorula</u> and/or an enzyme obtained from any of these microorganisms.

Preliminary Amendment

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4. (original): The process according to claim 2, wherein the enzyme source is a cultured product of a microorganism that has the activity of selectively reducing the compound represented by said formula (1) to produce the S-isomer and that belongs to genus Rhodococcus or genus Rhodotorula and/or an enzyme obtained from any of these microorganisms.

- 5. (original): The process according to claim 4, wherein the enzyme source that selectively produces the S-isomer is a cultured product of <u>Rhodococcus</u> sp. or <u>Rhodotorula</u> <u>glutinis</u> and/or an enzyme obtained from any of these microorganisms.
- 6. (original): The process according to claim 4, wherein the enzyme source that selectively produces the S-isomer is a cultured product of <u>Escherichia coli</u> HB101 (pNTRS) (FERM BP-08545) or <u>Escherichia coli</u> HB101 (pNTRGG1) (FERM BP-7858) and/or an enzyme obtained from any of these microorganisms.
- 7. (original): The process according to claim 2, wherein the enzyme source is a cultured product of a microorganism that has the activity of selectively reducing the compound represented by said formula (1) to produce the R-isomer and that belongs to genus <u>Candida</u> or genus <u>Devosia</u> and/or an enzyme obtained from any of these microorganisms.

8. (original): The process according to claim 7, wherein the enzyme source that selectively produces the R-isomer is a cultured product of <u>Candida malis</u>, <u>Candida magnoliae</u>, or <u>Devosia riboflavina</u> and/or an enzyme obtained from any of these microorganisms.

9. (original): The process according to claim 7, wherein the enzyme source that selectively produces the R-isomer is a cultured product of <u>Escherichia coli</u> HB101 (pNTS1G) (FERM BP-5835), <u>Escherichia coli</u> HB101 (pNTFPG) (FERM BP-7117), or <u>Escherichia coli</u> HB101 (pNTDRG1) (FERM BP-08458) and/or an enzyme obtained from any of these microorganisms.

10. (currently amended): The process according to <u>claim 1 any one of claims 1 to 9</u>, wherein 5-hydroxy-2-pentanone represented by said formula (1) produced by hydrolyzing 2-acetyl-γ-butyrolactone represented by formula (5):

in the presence of an acid is used as a starting material.

11. (currently amended): A process for producing an optically active 1-substituted 2-methylpyrrolidine represented by formula (4):

$$N_{R^2}$$
 (4)

(wherein R² represents a hydrogen atom, a hydroxyl group, a methoxy group, a benzyloxy group, a substituted or unsubstituted alkyl group having 1 to 12 carbon atoms, a substituted or unsubstituted aralkyl group having 7 to 12 carbon atoms, or a substituted or unsubstituted aryl group having 6 to 12 carbon atoms, and * represents an asymmetric carbon atom) comprising sulfonylating the optically active 1,4-pentanediol represented by said formula (2) produced by the processary one of the processes according to claim 1 claims 1 to 10 to convert it to an optically active disulfonate compound represented by formula (3):

$$SO_2R^1$$
 SO_2R^1
 SO_2R^1
 SO_2R^1

(wherein R¹ represents a substituted or unsubstituted alkyl group having 1 to 12 carbon atoms, a substituted or unsubstituted aralkyl group having 7 to 12 carbon atoms, or a substituted or unsubstituted aryl group having 6 to 12 carbon atoms, and * represents an asymmetric carbon atom), and reacting the compound with an amine.

12. (original): The process according to claim 11, wherein R¹ is a methyl group or a 4-methyphenyl group and R² is a benzyl group.